## In the claims

## 1-45 (canceled).

- 46 (currently amended). A method for treating a fibrotic disease comprising administering to a patient having a fibrotic disease a therapeutically effective amount of a composition comprising a pharmaceutically acceptable carrier and a polypeptide comprising SEQ ID NO: 2 or a salt thereof, wherein said fibrotic disease is lung fibrosis or liver fibrosis.
- 47 (previously presented). The method according to claim 46, wherein the fibrotic disease is lung fibrosis.
- 48 (previously presented). The method according to claim 46, wherein the polypeptide is glycosylated at one or more sites.
- 49 (previously presented). The method according to claim 46, wherein the polypeptide comprising SEQ ID NO: 2 is a fusion protein.
- 50 (previously presented). The method according to claim 49, wherein the fusion protein comprises an immunoglobulin Fc region fused to SEQ ID NO: 2.
  - 51-54 (canceled).
- 55 (previously presented). The method according to claim 46, wherein the polypeptide consists of SEQ ID NO: 2.
  - 56 (canceled).

- 57 (previously presented). The method according to claim 46, wherein the composition further comprises an interferon.
- 58 (previously presented). The method according to claim 57, wherein the interferon is interferon-B.
- 59 (previously presented). The method according to claim 46, wherein a composition comprising an interferon is administered to said patient simultaneously, sequentially, or separately with a composition comprising a pharmaceutically acceptable carrier and SEQ ID NO: 2.
- 60 (previously presented). The method according to claim 46, wherein said fibrotic disease is liver fibrosis
- 61 (new). The method according to claim 46, wherein said composition comprises a pharmaceutically acceptable carrier and a polypeptide comprising SEQ ID NO: 2.
- 62 (new). The method according to claim 46, wherein said composition comprises a pharmaceutically acceptable carrier and a salt of a polypeptide comprising SEQ ID NO: 2.
- 63 (new). The method according to claim 62, wherein said salt is a sodium, calcium, ammonium, ferric or zinc salt.
- 64 (new). The method according to claim 62, wherein said salt is a triethanolamine, arginine, lysine, piperidine or procaine salt.
  - 65 (new). The method according to claim 62, wherein said salt is an acid addition salt.
- 66 (new). The method according to claim 65, wherein said acid addition salt formed by the addition of hydrochloric, sulfuric, acetic or oxalic acid.